

Part I: Description of consignment	I.1. Consignor		I.2. IMSOC reference		I.2.a. Local reference	
	Name				I.3. Central Competent Authority	
	Address				I.4. Local Competent Authority	
	Country		ISO Code			
	I.5. Consignee			I.6. Operator conducting assembly operations independently of an establishment		
	Name			Name		
	Address			Address		
	Country			Approval Number		
				Country		
				ISO Code		
I.7. Country of origin			ISO Code		I.9. Country of destination	
					ISO Code	
I.8. Region of origin			Code		I.10. Region of destination	
					Code	
I.11. Place of dispatch			I.12. Place of destination			
Name			Name			
Address			Address			
Approval Number			Approval Number			
Country			Country			
			ISO Code			
I.13. Place of loading			I.14. Date and time of departure			
Name						
Address						
Approval Number						
Country			ISO Code			
I.15. Means of Transport			I.16. Transporter			
Mode	International transport document	Identification	Name			
			Address			
			Activity ID			
			Country			
			ISO Code			
			I.17. Accompanying documents			
			Commercial document reference			
			Date of issue			
			Country			
			Place of issue			
I.18. Transport conditions						
Frozen <input type="checkbox"/>		Chilled <input type="checkbox"/>		Ambient <input type="checkbox"/>		
I.19. Container No / Seal No						
I.20. Certified as Germinal products <input type="checkbox"/>						
I.21. For transit through a third country <input type="checkbox"/>						
Third country		ISO Code				
Exit point		BCP code				
Entry point		BCP code				
I.22. For transit through Member State(s) <input type="checkbox"/>			I.23. For export <input type="checkbox"/>			
Member State			Third country			
ISO Code			ISO Code			
			Exit point			
			BCP code			
I.25. Journey Log						
I.26. Total number of packages		I.27. Total quantity		I.28. Total gross weight		
I.30. Description of consignment						
Commodity	Species	Identification Number	Quantity	Nature of commodity		
Identification Mark	Package count	Date of collection / production	Plant / Establishment / Centre	Type		

Part II: Certification	II. Health information								
	<p>I, the undersigned official veterinarian, hereby certify that:</p> <p>II.1. The ova/embryos(1) described in Part I:</p> <p style="margin-left: 40px;">II.1.1. were collected, processed and stored under conditions which meet the requirements of Directive 92/65/EEC;</p> <p style="margin-left: 40px;">II.1.2. come from donor female swine which meet the requirements of Chapter IV of Annex D to Directive 92/65/EEC;</p> <p style="margin-left: 40px;">II.1.3. meet the requirements of Chapter III of Annex D to Directive 92/65/EEC.</p> <p><input type="checkbox"/> either [II.2. In the case of embryos,</p> <p style="margin-left: 40px;">II.2.1. the semen used for fertilisation meets the requirements of Directive 90/429/EEC;</p> <p style="margin-left: 40px;">II.2.2. the embryos have been washed with trypsin(2).]</p> <p>(1) <input type="checkbox"/> or [II.2. In the case of ova, the ova comes from a donor female swine which meets the conditions of Article 1 of Decision 2008/185/EC(2).]</p>								
<p>Notes</p> <p>This animal health certificate shall be completed according to the notes for the completion of certificates provided for in Chapter 2 of Annex I to Commission Implementing Regulation (EU) 2020/2235.</p> <p>Part I:</p> <p>Box I.11: Place of destination shall correspond to the embryo collection team of oocytes/embryos collection.</p> <p>Box I.12: Place of destination shall correspond to the embryo collection team, embryo production team, germinal product processing establishment, germinal product storage centre or to the establishment of oocytes/embryos destination.</p> <p>Box I.19: Identification of container and Seal number shall be indicated.</p> <p>Box I.30: "Type": specify if: in vivo derived embryos, in vivo derived oocytes, in vitro produced embryos or micromanipulated embryos.</p> <p style="margin-left: 40px;">Identification number shall correspond to the official identification of the animal.</p> <p style="margin-left: 40px;">Date of collection shall be indicated in the following format: dd/mm/yyyy.</p> <p style="margin-left: 40px;">Approval number of the team shall correspond to the embryo collection team of oocytes/embryos collection indicated in Box I.11.</p> <p>Part II:</p> <p>(1) Delete as appropriate.</p> <p>(2) This condition applies only to ova and embryos which originate in the Member States or regions thereof not listed in Annexes I and II to Decision 2008/185/EC (OJ L 59, 4.3.2008, p. 19) and destined to the Member States or regions thereof so listed. It shall also apply to movements from Member States or regions thereof listed in Annex II of Decision 2008/185/EC to Member States or regions thereof listed in Annex I of Decision 2008/185/EC.</p>									
<p>Certifying Officer/Official veterinarian</p> <table style="width: 100%; border: none;"> <tr> <td style="width: 50%; border: none;">Name (in capital letters)</td> <td style="width: 50%; border: none;">Authority name</td> </tr> <tr> <td style="border: none;">Date of signature</td> <td style="border: none;">Signature</td> </tr> <tr> <td style="border: none;">Stamp</td> <td style="border: none;"></td> </tr> </table>				Name (in capital letters)	Authority name	Date of signature	Signature	Stamp	
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